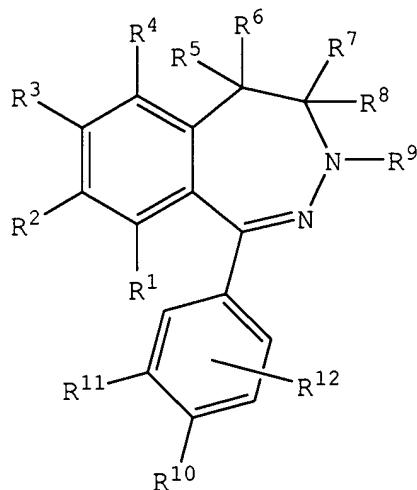


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This listing of the claims will replace all prior versions and listings of claims in the application:

Listing of Claims:

Claim 1 (previously presented): A compound of Formula I:



wherein

R¹, R², R³ and R⁴ are independently
H,
HO,
R¹³O-,
R¹³S-,
halogen,
C1-C3-alkyl,
CF₃,
R¹⁴CO₂-,
R¹⁴O₂C-,
R¹⁴CO-
R¹⁴CONH-,

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$R^{14}NHCO-$,

$R^{14}NHCO_2-$,

$R^{14}OCONH-$,

$R^{14}O_2S-$,

$R^{14}OS-$, or

$R^{15}R^{16}N-$; or

R^1 and R^2 , or R^2 and R^3 , or R^3 and R^4 taken together can be

$-SCH_2S-$,

$-SCH_2O-$,

$-OCH_2S-$,

$-SCH_2CH_2S-$,

$-SCH_2CH_2O-$, or

$-OCH_2CH_2S-$;

wherein one of R^1 , R^3 and R^4 must be C1-C3-alkoxy or C1-C3-alkylthio group;

R^5 , R^6 , R^7 and R^8 are independently

H,

C1-C6-alkyl,

C3-C6-alkenyl,

C3-C6-cycloalkyl,

phenyl or substituted phenyl, wherein the phenyl is substituted with one or two substituents, C1-C3-alkyl, halogen, $R^{13}O-$, CF_3- , $R^{14}O_2S-$, $R^{14}OS-$, $R^{14}CO-$, $R^{14}CO_2-$, $R^{14}O_2C-$, $R^{14}CONH-$, $R^{14}NHCO-$; or

R^5 and R^6 taken together can be C3-C6-cycloalkyl;

R^7 and R^8 taken together can be C3-C6-cycloalkyl;

R^9 is

$R^{15}R^{16}NCO-$,

$R^{15}R^{16}NCS-$,

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$R^{17}OCO-$,
 $R^{15}CO-$,
 $R^{15}R^{16}NCH_2CO-$,
 $R^{14}O_2C-(CH_2)_n-$,
 $R^{15}R^{16}NCO-(CH_2)_n-$,
 $NC-(CH_2)_n-$,
H,
C1-C6-alkyl,
C3-C6-alkenyl, or
C3-C6-cycloalkyl; or

R^8 and R^9 taken together can be

$-(CH_2)_mCH_2(R^{15})NCO-$,
 $-(CH_2)_mCH_2OCO-$, or
 $-(CH_2)_mCH_2CH_2CO-$;

R^{10} and R^{11} are independently

H,
 $R^{15}R^{16}N-$,
 $R^{14}HNCO-$, or
 $R^{14}CONH-$;

R^{12} is

H,
halogen,
HO,
 $R^{13}O-$,
 $R^{15}R^{16}N-$,
C1-C3-alkyl,
 CF_3 ,
 $R^{14}CO_2-$,
 $R^{14}CO-$, or

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$R^{14}CONH-$;
 R^{13} is C1-C3-alkyl;
 R^{14} is H or C1-C3-alkyl;
 R^{15} and R^{16} are independently
H,
C1-C10-alkyl,
C1-C6-perfluoroalkyl,
C3-C10-alkenyl, or
C3-C6-cycloalkyl; or
 R^{15} and R^{16} taken together can be C3-C6-cycloalkyl;
 R^{17} is C1-C6-alkyl, C3-C6-alkenyl, or C3-C6-cycloalkyl;
n is 1 to 6;
m is 0 to 2;
and pharmaceutically acceptable salts thereof;
wherein R^{10} and R^{11} cannot be both H.

Claim 2 (previously presented): The compound of claim 1 of Formula I wherein one of the substituents of R^1 , R^3 and R^4 must be C1-C3-alkylthio group or C1-C3-alkoxy group, the other substituents are independently H, $R^{13}O-$, $R^{13}S-$, halogen, or C1-C3-alkyl;

R^2 and R^3 taken together can be $-SCH_2S-$, SCH_2O- , or $-OCH_2S-$;
 R^9 is
 $R^{15}R^{16}NCO-$,
 $R^{15}R^{16}NCS-$,
 $R^{17}OCO-$,
 $R^{15}CO-$, or
H;

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R^{10} and R^{11} are independently H, H_2N- , or CH_3CONH- ;
and pharmaceutically acceptable salts thereof.

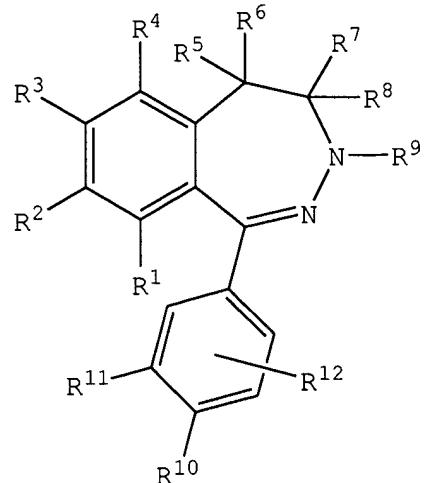
Claim 3 (previously presented): A composition comprising the compound of claim 2 and a pharmaceutically acceptable carrier.

Claims 4-7 (canceled).

Claim 8 (previously presented): A composition comprising the compound of claim 1 and a pharmaceutically acceptable carrier.

Claim 9 (canceled).

Claim 10 (currently amended): A method for treating a patient suffering from ischemia, epilepsy or stroke, the method comprising administering to the patient, in an effective amount to alleviate the symptoms of the ischemia, epilepsy or stroke, a compound of Formula I:



wherein

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R^1 , R^2 , R^3 and R^4 are independently

H,

HO,

$R^{13}O^-$,

$R^{13}S^-$,

halogen,

C1-C3-alkyl,

CF_3 ,

$R^{14}CO_2^-$,

$R^{14}O_2C^-$,

$R^{14}CO^-$,

$R^{14}CONH^-$,

$R^{14}NHCO^-$,

$R^{14}NHCO_2^-$,

$R^{14}OCONH^-$,

$R^{14}O_2S^-$,

$R^{14}OS^-$, or

$R^{15}R^{16}N^-$; or

R^1 and R^2 , or R^2 and R^3 , or R^3 and R^4 taken together can be

$-SCH_2S^-$,

$-SCH_2O^-$,

$-OCH_2S^-$,

$-SCH_2CH_2S^-$,

$-SCH_2CH_2O^-$, or

$-OCH_2CH_2S^-$;

wherein one of R^1 , R^3 and R^4 must be C1-C3-alkoxy or C1-C3-alkylthio group;

R^5 , R^6 , R^7 and R^8 are independently

H,

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C1-C6-alkyl,

C3-C6-alkenyl,

C3-C6-cycloalkyl,

phenyl or substituted phenyl, wherein the phenyl is substituted with one or two substituents, C1-C3-alkyl, halogen, $R^{13}O^-$, CF_3^- , $R^{14}O_2S^-$, $R^{14}OS^-$, $R^{14}CO^-$, $R^{14}CO_2^-$, $R^{14}O_2C^-$, $R^{14}CONH^-$, $R^{14}NHCO^-$; or

R^5 and R^6 taken together can be C3-C6-cycloalkyl;

R^7 and R^8 taken together can be C3-C6-cycloalkyl;

R^9 is

$R^{15}R^{16}NCO^-$,

$R^{15}R^{16}NCS^-$,

$R^{17}OCO^-$,

$R^{15}CO^-$,

$R^{15}R^{16}NCH_2CO^-$,

$R^{14}O_2C-(CH_2)_n^-$,

$R^{15}R^{16}NCO-(CH_2)_n^-$,

$NC-(CH_2)_n^-$,

H,

C1-C6-alkyl,

C3-C6-alkenyl, or

C3-C6-cycloalkyl; or

R^8 and R^9 taken together can be

$-(CH_2)_mCH_2(R^{15})NCO^-$,

$-(CH_2)_mCH_2OCO^-$, or

$-(CH_2)_mCH_2CH_2CO-[,]$;

R^{10} and R^{11} are independently

H,

$R^{15}R^{16}N^-$,

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$R^{14}HNCO-$, or
 $R^{14}CONH-$;
 R^{12} is
H,
halogen,
HO,
 $R^{13}O-$,
 $R^{15}R^{16}N-$,
C1-C3-alkyl,
 CF_3 ,
 $R^{14}CO_2-$,
 $R^{14}CO-$, or
 $R^{14}CONH-$;
 R^{13} is C1-C3-alkyl;
 R^{14} is H or C1-C3-alkyl;
 R^{15} and R^{16} are independently
H,
C1-C10-alkyl,
C1-C6-perfluoroalkyl,
C3-C10-alkenyl, or
C3-C6-cycloalkyl; or
 R^{15} and R^{16} taken together can be C3-C6-cycloalkyl;
 R^{17} is C1-C6-alkyl, C3-C6-alkenyl, or C3-C6-cycloalkyl;
n is 1 to 6;
m is 0 to 2;
and pharmaceutically acceptable salts thereof;
wherein R^{10} and R^{11} cannot be both H,
in combination with a pharmaceutically acceptable carrier.

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Claim 11 (previously presented): The method of claim 10 wherein, in the compound of Formula I,

one of the substituents of R¹, R³ and R⁴ must be C1-C3-alkylthio group or C1-C3-alkoxy group, the other substituents are independently H, R¹³O-, R¹³S-, halogen, or C1-C3-alkyl;

R² and R³ taken together can be -SCH₂S-, -SCH₂O-, or -OCH₂S-;

R⁹ is

R¹⁵R¹⁶NCO-,

R¹⁵R¹⁶NCS-,

R¹⁷OCO-,

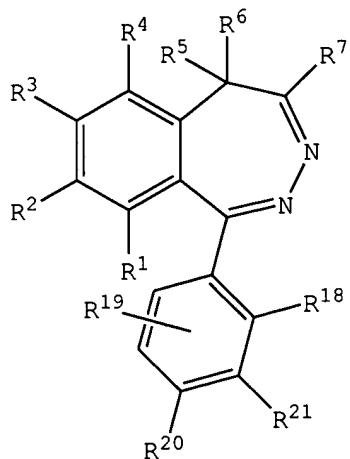
R¹⁵CO-, or

H;

R¹⁰ and R¹¹ are independently H, H₂N-, or CH₃CONH-; and pharmaceutically acceptable salts thereof.

Claim 12-15 (canceled).

Claim 16 (currently amended): A compound of Formula II:



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wherein

R^1 and R^4 are independently

H,
HO,
 $R^{13}O_-$,
 $R^{13}S_-$,
halogen,
C1-C3-alkyl,
 CF_3 ,
 $R^{14}CO_2-$,
 $R^{14}O_2C-$,
 $R^{14}CO_-$,
 $R^{14}CONH-$,
 $R^{14}NHCO-$,
 $R^{14}NHCO_2-$,
 $R^{14}OCONH-$,
 $R^{14}O_2S_-$,
 $R^{14}OS-$, or
 $R^{15}R^{16}N-$; or

R^2 is one of H, HO, $R^{13}O_-$, halogen, C1-C3-alkyl, CF_3 , $R^{14}CO_2-$, $R^{14}O_2C-$, $R^{14}CO_-$, $R^{14}CONH-$, $R^{14}NHCO-$, $R^{14}NHCO_2-$, $R^{14}OCONH-$, $R^{14}O_2S_-$, $R^{14}OS-$, $R^{13}S_-$ and $R^{15}R^{16}N-$ when R^3 is one of HO, halogen, C1-C3-alkyl, CF_3 , $R^{14}CO_2-$, $R^{14}O_2C-$, $R^{14}CO_-$, $R^{14}CONH-$, $R^{14}NHCO-$, $R^{14}NHCO_2-$, $R^{14}OCONH-$, $R^{14}O_2S_-$, $R^{14}OS-$, $R^{13}S_-$ and $R^{15}R^{16}N-$; or

R^2 is one of H, HO, halogen, C1-C3-alkyl, CF_3 , $R^{14}CO_2-$, $R^{14}O_2C-$, $R^{14}CO_-$, $R^{14}CONH-$, $R^{14}NHCO-$, $R^{14}NHCO_2-$, $R^{14}OCONH-$, $R^{14}O_2S_-$, $R^{14}OS-$, $R^{13}S_-$ and $R^{15}R^{16}N-$ when R^3 is one of H, HO, $R^{13}O_-$, halogen, C1-C3-alkyl, CF_3 , $R^{14}CO_2-$, $R^{14}O_2C-$, $R^{14}CO_-$, $R^{14}CONH-$, $R^{14}NHCO-$, $R^{14}NHCO_2-$, $R^{14}OCONH-$, $R^{14}O_2S_-$, $R^{14}OS-$, $R^{13}S_-$ and $R^{15}R^{16}N-$; or

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R^1 and R^2 , or R^2 and R^3 , or R^3 and R^4 taken together can be
- SCH_2S- ,
- SCH_2O- ,
- OCH_2S- ,
- SCH_2CH_2S- ,
- SCH_2CH_2O- , or
- OCH_2CH_2S- ;

wherein one of the substituents of R^1 , R^3 and R^4 must be C1-C3-alkoxy or C1-C3-alkylthio group;

R^5 , R^6 and R^7 are independently

H,
C1-C6-alkyl,
C3-C6-alkenyl,
C3-C6-cycloalkyl,

phenyl or substituted phenyl, wherein the phenyl is substituted with one or two substituents, C1-C3-alkyl, halogen, $R^{13}O^-$, CF_3^- , $R^{14}O_2S^-$, $R^{14}OS^-$, $R^{14}CO^-$, $R^{14}CO_2^-$, $R^{14}O_2C^-$, $R^{14}CONH^-$, $R^{14}NHCO^-$; or

R^5 and R^6 taken together can be C3-C6-cycloalkyl;

R^{13} is C1-C3-alkyl;

R^{14} is H or C1-C3-alkyl;

R^{15} and R^{16} are independently

H,
C1-C10-alkyl,
C1-C6-perfluoroalkyl,
C3-C10-alkenyl, or
C3-C6-cycloalkyl; or

R^{15} and R^{16} taken together can be C3-C6-cycloalkyl;

R^{18} and R^{19} are independently

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H,
halogen,
C1-C3-alkyl,
 $R^{14}O^-$,
 CF_3^- , or
 $R^{14}CO_2^-$;

R^{20} and R^{21} are independently

H,
 $R^{15}R^{16}N^-$,
 $R^{15}HNC(NH)^-$ or
 $R^{14}CONH^-$;

and pharmaceutically acceptable salts thereof;

wherein R^{20} and R^{21} cannot both be H.

Claim 17 (previously presented): The compound of claim 16 of Formula II

wherein one of the substituents of R^1 , R^3 and R^4 must be C1-C3-alkylthio group or C1-C3-alkoxy group, the other substituents are independently H, $R^{13}O^-$, $R^{13}S^-$, halogen, or C1-C3-alkyl;

R^2 and R^3 taken together can be $-SCH_2S^-$, $-SCH_2O^-$, or $-OCH_2S^-$;

R^{20} and R^{21} are independently H, H_2N^- , or CH_3CONH^- ;

and pharmaceutically acceptable salts thereof.

Claim 18 (previously presented): A composition comprising the compound of claim 17 and a pharmaceutically acceptable carrier.

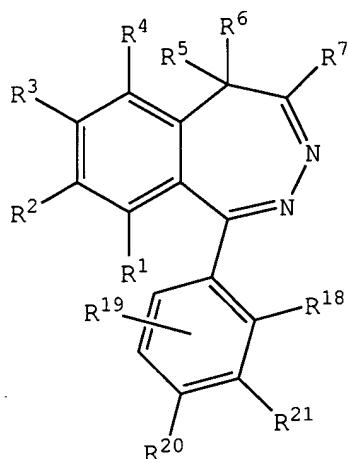
Claims 19-22 (canceled).

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Claim 23 (previously presented): A composition comprising the compound of claim 16 and a pharmaceutically acceptable carrier.

Claim 24 (canceled).

Claim 25 (currently amended): A method for treating a patient, the method comprising administering to the patient, in an effective amount to alleviate the symptoms of the ischemia, epilepsy or stroke, a compound of Formula II:



wherein

R¹ and R⁴ are independently

H,
HO,
R¹³O-,
R¹³S-,
halogen,
C1-C3-alkyl,
CF₃,
R¹⁴CO₂-,

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$R^{14}O_2C-$,
 $R^{14}CO-$,
 $R^{14}CONH-$,
 $R^{14}NHCO-$,
 $R^{14}NHCO_2-$,
 $R^{14}OCONH-$,
 $R^{14}O_2S-$,
 $R^{14}OS-$, or
 $R^{15}R^{16}N-$; or

R^2 is one of H, HO , $R^{13}O_-$, halogen, C1-C3-alkyl, CF_3 , $R^{14}CO_2-$, $R^{14}O_2C-$, $R^{14}CO-$, $R^{14}CONH-$, $R^{14}NHCO-$, $R^{14}NHCO_2-$, $R^{14}OCONH-$, $R^{14}O_2S-$, $R^{14}OS-$ and $R^{15}R^{16}N-$ when R^3 is one of HO, halogen, C1-C3-alkyl, CF_3 , $R^{14}CO_2-$, $R^{14}O_2C-$, $R^{14}CO-$, $R^{14}CONH-$, $R^{14}NHCO-$, $R^{14}NHCO_2-$, $R^{14}OCONH-$, $R^{14}O_2S-$, $R^{14}OS-$, $R^{13}S-$ and $R^{15}R^{16}N-$; or

R^2 is one of H, HO, halogen, C1-C3-alkyl, CF_3 , $R^{14}CO_2-$, $R^{14}O_2C-$, $R^{14}CO-$, $R^{14}CONH-$, $R^{14}NHCO-$, $R^{14}NHCO_2-$, $R^{14}OCONH-$, $R^{14}O_2S-$, $R^{14}OS-$ and $R^{15}R^{16}N-$ when R^3 is one of H, HO, $R^{13}O_-$, halogen, C1-C3-alkyl, CF_3 , $R^{14}CO_2-$, $R^{14}O_2C-$, $R^{14}CO-$, $R^{14}CONH-$, $R^{14}NHCO-$, $R^{14}NHCO_2-$, $R^{14}OCONH-$, $R^{14}O_2S-$, $R^{14}OS-$, $R^{13}S-$ and $R^{15}R^{16}N-$; or

R^1 and R^2 , or R^2 and R^3 , or R^3 and R^4 taken together can be

$-SCH_2S-$,
 $-SCH_2O-$,
 $-OCH_2S-$,
 $-SCH_2CH_2S-$,
 $-SCH_2CH_2O-$, or
 $-OCH_2CH_2S-$;

wherein one of the substituents of R^1 , R^3 and R^4 must be C1-C3-alkoxy or C1-C3-alkylthio group;

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R^5 , R^6 , and R^7 are independently
H,
C1-C6-alkyl,
C3-C6-alkenyl,
C3-C6-cycloalkyl,
phenyl or substituted phenyl, wherein the phenyl is
substituted with one or two substituents, C1-C3-alkyl, halogen,
 $R^{13}O^-$, CF_3^- , $R^{14}O_2S^-$, $R^{14}OS^-$, $R^{14}CO^-$, $R^{14}CO_2^-$,
 $R^{14}O_2C^-$, $R^{14}CONH^-$, $R^{14}NHCO^-$; or
 R^5 and R^6 taken together can be C3-C6-cycloalkyl;
 R^{13} is C1-C3-alkyl;
 R^{14} is H or C1-C3-alkyl;
 R^{15} and R^{16} are independently
H,
C1-C10-alkyl,
C1-C6-perfluoroalkyl,
C3-C10-alkenyl, or
C3-C6-cycloalkyl; or
 R^{15} and R^{16} taken together can be C3-C6-cycloalkyl;
 R^{18} and R^{19} are independently
H,
halogen,
C1-C3-alkyl,
 $R^{14}O^-$,
 CF_3^- , or
 $R^{14}CO_2^-$;
 R^{20} and R^{21} are independently
H,
 $R^{15}R^{16}N^-$,

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$R^{15}HNC(NH) -$ or

$R^{14}CONH -$;

and pharmaceutically acceptable salts thereof;

wherein R^{20} and R^{21} cannot both be H[.]

in combination with a pharmaceutically acceptable carrier.

Claim 26 (previously presented): The method of claim 25
wherein, in the compound of Formula II

wherein one of the substituents of R^1 , R^3 and R^4 must be Cl-
C3-alkylthio group or Cl-C3-alkoxy group, the other substituents
are independently H, $R^{13}O -$, $R^{13}S -$, halogen, or Cl-C3-alkyl;

R_2 and R_3 taken together can be $-SCH_2S -$, $-SCH_2O -$, or $-OCH_2S -$;

R^{20} and R^{21} are independently H, $H_2N -$, or $CH_3CONH -$;

and pharmaceutically acceptable salts thereof.

Claims 27-30 (canceled).